

## **Product Data Sheet**

## Anti-IL-17 Antibody

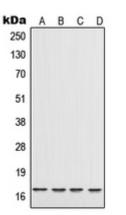
Catalog #	Source	Reactivity	Applications	
CPA1598	Rabbit	-	WB, IH	
			·	
Description		Rabbit polyclonal antibody to	) IL-17	
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within the center	
		region of human IL-17. The e	xact sequence is proprietary.	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	s of IL-17 protein.	
Clonality Polyclonal				
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/100)	
Gene Symbol		IL17A		
Alternative Na	ames	CTLA8; IL17; Interleukin-17A	IL-17; IL-17A; Cytotoxic T-lymphocyte-associated	
		antigen 8; CTLA-8		
Entrez Gene		3605 (Human); 16171 (Mous	e)	
SwissProt		Q16552 (Human); Q62386 (N	1ouse); Q61453 (Rat)	
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
		freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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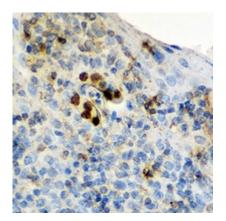




For research purposes only, not for human use

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Western blot analysis of IL-17 expression in A20 (A), THP1 (B), Raw264.7 (C), H9C2 (D) whole cell lysates. (Predicted band size: 17 kD; Observed band size: 17 kD)



Immunohistochemical analysis of IL-17 staining in human tonsil formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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